Platform switching versus platform matching:

two-year results from a prospective randomized-controlled multicenter study

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Background and Aim:

Platform switching concept is based on the discrepancy between the prosthetic abutments of a smaller diameter in relation to the implant platform diameter and clinically seems to preserve crestal bone height and soft tissue levels increasing the quality outcomes in treatments with dental implants and the patient satisfaction. However, it is well-known that the lack of well-designed prospective randomized clinical trials evaluating the efficiency of platform switching (PS) versus platform matching (PM) placed in partially edentulous mandibles.

The purpose of this five-year prospective randomized multicenter study was to assess the differences in bone level changes between CAMLOG® SCREW-LINE implants supporting single crowns in the posterior mandible restored either with platform matching or platform switching abutments (FDI positions 37-34 and/or 44-47). The secondary objectives included implant success (Buser et al. 2002) and survival rate, performance of the restorative components, nature and frequency of the adverse events.

This paper presents interim results obtained in up to two years.

Material and methods:

The prospective multicenter randomized clinical study was performed in three centers located in Germany (two) and Portugal (one). The study was approved by the competent Ethics Committees (FECI 09/1308 and CES/0156) and performed in accordance with the Declaration of Helsinki (2008).

Patients ≥18 years old missing two or more adjacent teeth in the posterior mandible and with a natural tooth mesially to the most proximal implant site.

Free end situation was allowed and opposing dentition must be natural teeth or implant supported fixed restorations.

Following implant placement (i.e. before placing the healing cap) patients were randomized either in the group of abutment for PM restoration or in the group for PS. All patients signed the detailed informed consent form before surgery.

Exclusion criteria

Individuals who presented uncontrolled systemic diseases or took medications interfering with bone metabolism or presenting abuse of drugs or alcohol, use of tobacco equivalent to >10 cigarettes/day or presenting handicaps that would interfere with the ability to perform adequate oral hygiene.

Material

-CAMLOG® SCREW-LINE Implants, Promote® plus:

- Diameter 3.8, 4.3 and 5 mm
- Length 9, 11 and 13 mm
- -Platform switching and platform matching prosthetic components.



Study status

Center	Patients	Implants Intent to treat	Implants per protocol	Comments	
Kiel	14	28	24 PS 12, Std 12	2 patients (4 implants) off-protocol	
Mainz	21	57	53 PS 29, Std 24	4 implants in two pat. off-protocol 3 patients with split-mouth design	
Coimbra	35	86	86 PS 42, Std 44	5 patients with split-mouth design	
TOTAL	70	167	163 PS 83 Std 80		

	Total patients / implants	Split-mouth pat.	Status
Randomized (In-protocol)	68 / 163	8	Completed
Loading*	67 / 160	8	Completed
12-month FU	67 / 160	8	Completed
24-month FU	61 / 144**	7	Ongoing

- * 3 implants non-osseointegrated (NOI) and explanted prior to loading (two lost, one mobile).
- ** 5 patients (14 implants) 24-month FU information pending; 1 patient with 2 implants died 20-month post-loading.

Demography Male / Female 22 / 17 20 /17 MEAN Age at surgery 53.29 (SD 10.44) 50.17 (SD 14.33) Smoking status 25 No smoker Former smoker 4 Current smoker 8

Implants description Implant position by randomization Implants by length and diameter in mm

Clinical Protocol Patient 304 Baseline Loading and

Prosthesis Delivery 10 ± 2 weeks after Surgery

Follow-up Visits

Standardized Radiographs







Initial

Statistical methods

The distance from the implant shoulder to the first crestal bone contact, at the mesial and distal side, was measured with standardized radiographs and averaged to represent the change of bone level over time per implant. Two-way ANOVA considering Center and Randomization as factors was used to evaluate the mean differences in bone level change at a significance level of 0.05.

Survival analysis was applied to calculate implant success and survival rate.

Results:

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ı	Bolle gail = positive value (+) Bolle loss = negative value (-)							
	Randomization	Platform switching		Platform matching		2-way ANOVA		
		N	MEAN ± SD	N	MEAN ± SD	Center effect	Randomization effect	
	Surgery to loading	76	- 0.53± 0.45	70	- 0.63 ± 0.70	Yes (p<0.01)	No (p=0.087)+	
	Loading – 12M*	76	0.10 ± 0.41	68	- 0.09 ± 0.50	Yes (p=0.024)	Yes (p=0.03)**	
	Loading – 24M**	69	0.26 ± 0.44	64	- 0.16 ± 0.65	Yes (p=0.027)	Yes (p<0.01)**	

* Despite statistically significant for centers 1 (0.046) and 2 (0.003) ** Overall difference between PS and PM attributable only to center 3 (p<0.01)

Bone level change two years post-loading per implant subdivided in 0.2 mm intervals and by randomization The mean bone level changes at two years post loading. Number of implants * 15.0 0.2 subdivided in mm intervals. In 81% of the implants in PS group and 48%in PM group bone gain was observed. A bone gain higher than 0.4 mm was observed in 33% of the implants with PS and only in 7,8% in PM group.

The survival rate of this study are 97.6% .in the PS group and 98.8% in the PM group (not statistically significant).

Conclusions:

At two-year post-loading the implants restored with platform switching abutments appear to preserve the crestal bone more predictably than the implants restored with platform match.

References:

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